National Phase of PCT/DK2003/000463

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E. Steiness

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## AMENDMENTS TO THE SPECIFICATION

## In the specification:

Please enter the enclosed sequence listing on page 43 of the specification and renumber the pages with the Claims and Abstract accordingly. In addition to the diskette, a paper copy of the Sequence Listing is provided herewith.

Please amend the specification as shown:

Please delete the paragraph on page 2, lines 1-7, and replace it with the following paragraph:

--Many molecular derivatives of GLP-1 have been reported. For instance, GLP-1 (7-37) is known. A variety of analogs have also been described eg., Gln<sup>9</sup>-GLP-1 (7-37) (SEQ ID NO: 15), D-Gln<sup>9</sup>-GLP-1 (7-37), acetyl-Lys<sup>9</sup>-GLP-1 (7-37) (SEQ ID NO: 16), Thr<sup>16</sup>-Lys<sup>18</sup>-GLP-1(7-37) (SEQ ID NO: 17), and Lys<sup>18</sup>-GLP-1 (7-37) (SEQ ID NO: 18), Gly<sup>8</sup>-GLP-1 (7-37) (SEQ ID NO: 19), Ser<sup>8</sup>-GLP-1 (7-37) (SEQ ID NO: 20). Other GLP-1 derivatives (sometimes called "variants") have also been reported particularly as acid addition salts, carboxylate salts, lower alkyl esters, and amides. See WO 91/11457 and Mojsov, S., *Int. J. Peptide Protein Research*, 40:333-343 (1992), and references cites therein.--

Please delete the paragraph on page 10, lines 13-16, and replace it with the following paragraph:

--The amino acid sequence of GLP-1 (7-37) is well-known and has the following sequence: NH<sub>2</sub>-His<sup>7</sup> -Ala-Glu-Gly<sup>10</sup> -Thr-Phe-Thr-Ser-Asp<sup>15</sup> -Val-Ser-Ser-Tyr-Leu<sup>20</sup> -Glu-Gly-Gln-Ala-Ala<sup>2520</sup> -Lys-Glu-Phe-Ile-Ala<sup>30</sup> -Trp-Leu-Val-Lys-Gly-Arg-Gly<sup>37</sup> -COOH (SEQ ID NO: <u>1</u>)--

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Please delete the paragraph on page 10, lines 18-25, and replace it with the following paragraph:

--A "GLP-1 analog" is defined as a molecule having a modification including one or more amino acid substitutions, deletions, inversions, or additions when compared with GLP-1. GLP-1 analogs include, for example, GLP-1 (7-34) (SEQ ID NO: 21) and GLP-1 (7-35) (SEQ ID NO: 22), GLP-1 (7-36) (SEQ ID NO: 23), Val<sup>8</sup>-GLP-1 (7-37) (SEQ ID NO: 24), Gln<sup>9</sup>-GLP-1 (7-37) (SEQ ID NO: 15), D-Gln<sup>9</sup>-GLP-1 (7-37), Thr<sup>16</sup>-Lys<sup>18</sup>-GLP-1(7-37) (SEQ ID NO: 17), and Lys<sup>18</sup>-GLP-1 (7-37) (SEQ ID NO: 18). Preferred GLP-1 analogs are GLP-1 (7-34) (SEQ ID NO: 21) and GLP (7-35) (SEQ ID NO: 22), which are disclosed in U.S. Pat. No. 5,118,666, and also GLP-1 (7-36) (SEQ ID NO: 23). These compounds are the biologically processed forms of GLP-1 having insulinotropic properties. Other GLP-1 analogs are disclosed in U.S. Pat. No. 5,545,618.--

Please delete the paragraph on page 17, line 16, to page 18, line 12, and replace it with the following paragraph:

--More particular examples of GLP-1 and GLP-1 related molecules including analogs thereof such as those disclosed in the PCT/DK00/00393 application. Such molecules include the following specific compounds:

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des Ser<sup>39</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Pro<sup>36</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Ala<sup>35</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Gly<sup>34</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Ser<sup>39</sup> -(Lys<sup>40</sup> (palmitoyl))exendin-4(1-39)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Gly<sup>34</sup> -(Lys<sup>40</sup> (palmitoyl))exendin-4(1-39)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Ala<sup>35</sup> -(Lys<sup>40</sup> (palmitoyl))exendin-4(1-39)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ ID NO:___),
des Pro<sup>36</sup> -(Lys<sup>40</sup> (palmitoyl))exendin-4(1-39)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ ID NO:___),
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Lys<sup>40</sup> (palmitoyl)exendin-4(1-39)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ-ID-NO:—),

des Pro<sup>36</sup>, Pro<sup>37</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub>.

Lys<sub>6</sub>-des Pro<sup>36</sup>, Pro<sup>37</sup>, Pro<sup>38</sup> -exendin-4(1-39)-NH<sub>2</sub>.

Asn(Glu)<sub>5</sub>-des Pro<sup>36</sup>, Pro<sup>37</sup>, Pro<sup>38</sup> -exendin-4(1-39)-NH<sub>2</sub>.

Lys<sub>6</sub>-des Pro<sup>36</sup>, Pro<sup>37</sup>, Pro<sup>38</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub>

Asn(Glu)<sub>5</sub>-des Pro<sup>36</sup>, Pro<sup>37</sup>, Pro<sup>38</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub>,

des Pro<sup>36</sup>, Pro<sup>37</sup>, Pro<sup>38</sup> -exendin-4(1-39)-Lys<sub>6</sub>-NH<sub>2</sub>.

Gly<sup>8</sup>-GLP-1 (7-36)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO: <u>6</u>)

Lys<sub>6</sub>-Gly<sup>8</sup>-GLP-1 (7-36)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO: 7),

Lys<sub>6</sub>-Gly<sup>8</sup>-GLP-1 (7-36)-NH<sub>2</sub> (SEQ ID NO: 8),

(Gly<sup>8</sup>,Lys<sup>37</sup>(palmitoyl)-GLP-1(7-36)(Human)-Lys<sub>7</sub>-NH<sub>2</sub> (SEQ ID NO: <u>9</u>),

(Gly<sup>8</sup>,Lys<sup>26</sup>(palmitoyl)-GLP-1(7-36)(Human)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO: 10),

(Gly<sup>8</sup>,Lys<sup>34</sup>(palmitoyl)-GLP-1(7-36)(Human)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO: 11),

Gly<sup>8</sup>-GLP-1 (7-36)-Lys<sub>8</sub>-NH<sub>2</sub> (SEQ ID NO: 12),

Gly<sup>8</sup>-GLP-1 (7-36)-Lys<sub>10</sub>-NH<sub>2</sub> (SEQ ID NO: 13),

Gly<sup>8</sup>-GLP-1 (7-37)-Lys<sub>6</sub>-NH<sub>2</sub> (SEQ ID NO: 14); and the free acid or pharmaceutically acceptable salt thereof.--

Please delete the paragraph on page 37, lines 12-19, and replace it with the following paragraph:

--Insulin standard for quantitative PCR. One μl first strand synthesis was used for PCR with the following insulin primers: 5'-AACCCACCCAGGCTTTTGTCA (SEQ ID NO: 2); 5'-CTTCCTCCCACGTCCAGTTGTTC-3 (SEQ ID NO: 3). The amplicon were inserted into the PCR 4-TOPO vector (invitrogen) and transformed into E.coli. The plasmids were purified and 2 μg of each were linearized with either Spe I or Not I. The linearized plasmids were *in-vitro* transcribed using T7 or T3 RNA polymerase. After *in-vitro* transcription, the template was removed by DNAse treatment. Subsequently, the mixture was phenol/chloroform-extracted and

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precipitated. After precipitation the RNA was dissolved to 1 mg/m1 in water.--

Please delete the paragraph on page 37, line 21, to page 38, line 3, and replace it with the following paragraph:

--Quantitative PCR. One μg of both standard and sample were subjected to first strand synthesis as described above. A dilution series of the insulin mRNA standard, together with the samples were subjected to quantitative PCR using the following probe (Mouse insulin Taqman probe, 110-138) and the above described primers:

5'-FAM-AGGCTCTCTACCTGGTGTGTGGGGAGCGT-Tamra-3' (SEQ ID NO: 4).

All PCR reactions were duplicates. The C<sub>t</sub> (threshold cycle) were measured and the initial concentration of insulin mRNA was calculated according to the standard curve.

Drugs: COMPOUND 1 (H-HGEGTFTSDLSKQMEEEAVRLFIEWLKNGGPSSGAP PSKKKKKK-NH2, (SEQ ID NO: 5) Batch: ZP15.65-3A) was produced at Zealand Pharma A/S using the Merifield technique.--